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Pubertal maturation and affective symptoms in adolescence and adulthood: evidence from a prospective birth cohort

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Short title: Puberty and life course affective symptoms

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

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Abstract

The higher prevalence of affective symptoms among women compared to men emerges in adolescence, and it has been associated with pubertal maturation. However, it remains unclear whether pubertal timing has long-term influences on affective symptoms. Using data from the British 1946 birth cohort, we investigated whether pubertal timing was associated with affective symptoms over the life course, distinguishing those with symptoms in adolescence only, symptoms in adulthood only, and symptoms in both adolescence and adulthood. In females, there was no evidence that early pubertal maturation was a risk factor for affective symptoms. However, those with particularly late menarche (≥ 15 years) showed a lower risk of adult-onset affective symptoms (OR = 0.54, 95% CI: 0.31, 0.95). This effect of late pubertal timing was not explained by a range of socio-behavioural factors. In contrast, in males, late pubertal timing was associated with increased risk of adolescent-onset affective symptoms that tracked into adulthood (OR = 2.10, 95% CI: 1.44, 3.06). This effect was partly explained by low pre-pubertal BMI. Sex-specific effects of pubertal timing on the long-term risk of affective symptoms might be due to different effects of gonadal hormonal on the CNS, as well as different social experiences during puberty.

Keywords: depression, anxiety, sex, adolescence, life course, puberty

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Depression and anxiety are among the most common forms of psychopathology in adolescence and adulthood (Costello, Mustillo, Erkanli, Keeler, & Angold, 2003). Recent estimates suggest that depression will become the second leading medical cause of disability in the world by 2020 (World Health Organization, 2001), and that the prevalence rate is rising among young people (Collishaw, Maughan, Goodman, & Pickles, 2004). It is known that adolescent emotional problems have long-term negative consequences across multiple domains (Richards & Abbott, 2009), maintain a chronic course for a significant proportion of youth affected (Colman, Wadsworth, Croudace, & Jones, 2007; Copeland, Shanahan, Costello, & Angold, 2011), and can increase risk for other mental and physical health problems (Copeland, Shanahan, Costello, & Angold, 2011; Gaysina, Hotopf, et al., 2011; Gaysina, Pierce, et al., 2011). Therefore, adolescence can be considered as a high-risk developmental period for the onset and intensification of depression and anxiety symptoms (Inderbitzen & Hope, 1995; Macaulay & Kleinknecht, 1989). Identification of pathways and processes through which negative outcomes are explained can provide information about modifiable targets for the development of intervention programs aimed at remediating risk outcomes among youth at risk.

Current developmentally sensitive etiologic models of affective disorders consider the potential role of pubertal maturation in development of psychopathology in adolescence (e.g., Angold & Costello, 2006). A growing body of evidence suggests that early pubertal maturation is associated with increased risk of depressive symptoms in girls, while the evidence in boys is limited and inconsistent (for review, see Reardon, Leen-Feldner, & Hayward, 2009). Moreover, very little attention has been paid to effects of pubertal timing on affective disorders across the life course. Therefore, it remains unclear whether the association between pubertal timing and affective symptoms in adolescence persists into adulthood, and what potential socio-behavioural mechanisms underlie this association. The

present research is therefore timely, as it addresses this gap in the literature by investigating associations between pubertal timing and affective symptoms across the life course using data from a prospective UK birth cohort.

Pubertal maturation and risk of affective disorders

Puberty can be considered as a “sensitive period” (Romeo, 2003), during which experiences exert particularly potent effects on brain development and related behavioural patterns (Andersen & Teicher, 2008). As such, the typical changes of puberty make it a high risk phase of the life course for vulnerable youth, rendering puberty a potentially important time for screening and intervention. Most studies of pubertal development and affective disorders focus on maturation *stage* or *timing*. Maturation stage refers to the degree of physical maturation at a given time point, often indexed by level of breast (in girls), genital (in boys), and pubic hair (in both boys and girls) development, and sometimes indexed by levels of pubertal hormones (e.g., Shirtcliff, Dahl, & Pollak, 2009). Timing describes the relative rank of an individual’s stage of development compared with same age and sex peers or the age at which an individual enters a particular stage of maturation (e.g., Marceau, Ram, Houts, Grimm, & Susman, 2011). Pubertal timing has been most commonly examined for association with affective symptoms (for review, see Reardon, Leen-Feldner, & Hayward, 2009). This measure is particularly valuable since it can be used to identify sub-groups of youth that might be at particular risk for affective disorders later in life.

Most, although not all, longitudinal studies following adolescent girls through puberty have found that early age at menarche was associated with increased risk of depressive symptoms during adolescence (Canals, Marti-Henneberg, Fernandez-Ballart, Cliville, & Domenech, 1992; Laitinen-Krispijn, van der, & Verhulst, 1999; Patton et al., 2008). There have been fewer studies in boys, and the evidence that exists is more inconsistent than that for girls, with some studies reporting a significant association between early pubertal maturation

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

and affective symptoms (Ge, Conger, & Elder, 2001), while others reporting an association between later pubertal maturation and depression (Angold, Costello, & Worthman, 1998; Conley & Rudolph, 2009; Laitinen-Krispijn, van der, & Verhulst, 1999), or no association (Graber, Seeley, Brooks-Gunn, & Lewinsohn, 2004; Patton, et al., 2008). The inconsistency in the results may be due to differences between studies in the measures of depression, anxiety, or combinations of both, and the small sample sizes of some studies meaning that they might lack statistical power to detect an association.

The prevalence of depression in adulthood is twice as high in women as in men (Angold & Costello, 2006). Since sex differences emerge around the age of puberty, it has been suggested that interactions between pubertal hormones and the adolescent brain may affect risk for psychopathology across the life course (Sisk & Zehr, 2005). However, few studies have considered pubertal timing in relation to affective disorders in adulthood (Graber, Seeley, Brooks-Gunn, & Lewinsohn, 2004; Harlow, Cohen, Otto, Spiegelman, & Cramer, 1999; Herva et al., 2004; Ryan, Carriere, Scali, Ritchie, & Ancelin, 2008), and the existing results are conflicting. One study found increased risk of current depressive symptoms in adult females to be associated with younger age at menarche (Harlow, Cohen, Otto, Spiegelman, & Cramer, 1999), while another found higher risk for depression in those with particularly late menarche (Herva, et al., 2004), and a third found no association (Ryan, Carriere, Scali, Ritchie, & Ancelin, 2008). The inconsistency in the findings may be explained by differences in age and/or ethnic origins of the samples of females (Hamlat, Stange, Abramson, & Alloy, 2014), as well as by differences in the measures of affective symptoms and pubertal timing used across the studies. Moreover, the existing studies did not distinguish groups of females with different age at onset of the symptoms (i.e. adolescent-onset versus adult-onset).

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

The evidence relating pubertal timing and later psychological distress in adult males is very limited. A study by Graber et al. (2004) observed that late-maturing men had elevated rates of lifetime history of disruptive behavior disorder and current substance use, but not depressive disorders, in young adulthood (age 24 years). In another study, Natsuaki, Biehl, and Ge (2009) demonstrated that that earlier maturing boys were at risk of manifesting the highest levels of depressed mood particularly around ages 15–16, but at older ages (up to age 23 years) the adverse effect of early maturation decreased.

Therefore it remains unclear whether differences in the timing of physical maturation have any long-term consequences for affective disorders in both men and women. It has been hypothesized that the negative impact of early puberty may not persist into adulthood (Dick, Rose, Viken, & Kaprio, 2000), or that those with later puberty can “catch up” and experience similar risk of mental health problems in adulthood to those with early or normative pubertal timing (Stattin & Magnusson, 1990).

Risk factors for pubertal maturation and affective disorders

There are several theories proposed to explain the observed associations between pubertal maturation and affective disorders, including biological (i.e., genetic, neuroendocrine) and socio-behavioural explanations (e.g., Angold, Costello, Erkanli, & Worthman, 1999; Costello, Mustillo, Erkanli, Keeler, & Angold, 2003; Ge & Natsuaki, 2009; Paikoff & Brooks-Gunn, 1991). It has been suggested that the association between pubertal maturation and affective symptoms might be explained by common environmental risk factors (Marceau et al., 2012). For example, adverse prenatal and early postnatal experiences may have long term consequences on growth and brain development that in turn can influence pubertal maturation and emotional health (dos Santos Silva et al., 2002; Colman, Ploubidis, Wadsworth, Jones, & Croudace, 2007). Risk factors can also mediate the association between pubertal maturation and affective disorders. Risk factors potentially

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

important for associations between pubertal maturation and affective symptoms in adolescence include perception of maturation status and change in social relationships (Cyranowski, Frank, Young, & Shear, 2000; Natsuaki et al., 2009). In addition to these, a range of other socio-behavioural factors need to be considered to explain the association with affective symptoms in later life. Early-maturing adolescents, and particularly girls, are at increased risk for unhealthy behaviours, including low levels of physical activity (Baker, Birch, Trost, & Davison, 2007; Lanza & Collins, 2002; van Jaarsveld, Fidler, Simon, & Wardle, 2007), and smoking (Jaszyna-Gasior et al., 2009; Lanza & Collins, 2002; van Jaarsveld, Fidler, Simon, & Wardle, 2007). Girls with early age at puberty are also at risk for higher levels of central adiposity (Hardy, Kuh, Whincup, & Wadsworth, 2006; Lakshman et al., 2008). They are more likely to start sexual practice earlier than normal developers and are at higher risk of teen pregnancy and early first childbearing (Talashek, Montgomery, Moran, Paskiewicz, & Jiang, 2000). These behaviors have also been associated with subsequent affective disorders (Chaiton, Cohen, O'Loughlin, & Rehm, 2009; Herva et al., 2006; Lancaster et al., 2010; Roshanaei-Moghaddam, Katon, & Russo, 2009). Early puberty is associated with lower school achievement and educational attainment in girls (Cavanagh, Riegle-Crumb, & Crosnoe, 2007), with lower educational attainment being associated with higher prevalence of mood and anxiety disorders and symptoms (Lorant et al., 2003). Therefore educational attainment may mediate the link between timing of pubertal maturation and affective symptoms.

Taken together, the existing evidence suggests that pubertal timing can be associated with affective psychopathology in adolescence and adulthood due to common risk factors; therefore, these factors need to be taken into account when studying the relations between pubertal maturation and affective psychopathology.

Present study

The present study examined the effects of pubertal timing on affective symptoms from adolescence to midlife using data from the Medical Research Council (MRC) National Survey of Health and Development (NSHD), a prospective birth cohort study originally of 5362 men and women born in Britain in 1946. Specifically, the study assessed whether any long-term association was dependent on the time at onset of affective symptoms and persistence of symptoms (i.e., those present only in adolescence, or only in adulthood, or in both adolescence and adulthood). The study also explored whether any observed association with adolescent-onset symptoms could be explained by common risk factors such as prenatal and pre-pubertal growth, or socioeconomic disadvantage; and whether any association with adult-onset symptoms could be explained by specific socio-behavioural mechanisms.

Method

Sample

The MRC NSHD is a birth cohort study of 2547 women and 2815 men born in Britain in one week in March 1946. There have been 22 follow-ups of the whole cohort between birth and age 53 years. The data collection received ethical approval from the North Thames MREC, and informed consent was given by respondents to each set of questions and measures. A total of 1972 men (69% of the original sample of men) and 1809 women (71% of the original sample of women) had information on affective symptoms across the life course and on pubertal timing at age 15 years.

Measures

Affective symptoms. At ages 13 and 15 years, teacher ratings were collected using a forerunner of the Rutter teacher questionnaire (Rutter, 1967). Teachers described aspects of the children's personality, behaviour, and mood on a 3-point scale. These questionnaires have

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

previously been subjected to classical linear factor analysis, with one factor comprising 11 items being identified as affective symptoms (depression and anxiety; Colman, Wadsworth, Croudace, & Jones, 2007; van Os, Jones, Lewis, Wadsworth, & Murray, 1997). This factor has been shown to be a good predictor of adult depression and anxiety (Colman, Wadsworth, Croudace, & Jones, 2007). Frequency and severity of affective symptoms were assessed in adulthood, with the Present State Examination (Wing, Cooper, & Sartorius, 1974) at 36 years, the Psychiatric Symptom Frequency scale (Lindelow, Hardy, & Rodgers, 1997) at 43 years, and the 28-item General Health Questionnaire (Goldberg & Hillier, 1979) at age 53 years.

Utilising these measures from adolescence to midlife (age 53 years), latent class analysis was previously employed to develop longitudinal profiles of affective symptoms (Colman, Ploubidis, Wadsworth, Jones, & Croudace, 2007). A single factor score representing affective symptoms at each of the five ages was estimated after confirmatory factor analysis. Due to the skewed distributions of these scores, a categorical variable with four groups (absence of symptoms, occasional symptoms, moderate symptoms and severe symptoms) was defined at each age. Using latent class analysis, six distinct profiles were identified ($N = 4627$): absence of symptoms (44.8% of sample), repeated moderate symptoms (33.6%), adult-onset moderate symptoms (11.3%), adolescent symptoms with good adult outcome (5.8%), adult-onset severe symptoms (2.9%) and repeated severe symptoms over the life course (1.7%). Because of our interest in the relationship between pubertal timing and onset and persistence of affective symptoms we defined four groups according to these two criteria: 1) absence of symptoms (44.8% of sample); 2) symptoms in adolescence only (5.8%); 3) symptoms in adulthood only (moderate and severe; 14.2%), 4) symptoms in both adolescence and adulthood (moderate and severe; 35.3%). Groups 2 and 4 represented adolescent-onset symptoms, whereas group 3 represented adult-onset symptoms. We have previously used this four-profile classification in order to test for association between

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

affective symptoms and BMI across the life course and have demonstrated that these four profiles have distinctive BMI trajectories in both men and women (Gaysina, Hotopf, et al., 2011).

Pubertal maturation. Age at menarche was used as the primary marker of pubertal timing for girls, and was obtained from mothers' reports at a medical examination by a school doctor at age 15 years. Five pubertal timing groups were defined based on age at menarche: ≤ 11.11 years, 12-12.11 years, 13-13.11 years, 14-14.11 years, and ≥ 15 years. Girls who had not reached menarche at the time of the examination were included in the latest group. In addition, other markers of pubertal maturation were collected during the examination: signs of breast development (yes or no); visible pigmented pubic hair (yes, profuse; yes, sparse; or no); visible axillary hair (yes or no). Information on presence of visible pigmented pubic hair and axillary hair was also collected for the boys, as were the development of genitalia (infantile; early; advanced) and voice breaking (no; starting; completely broken). Based on these observations, boys were classified as fully mature (advanced development of genitalia, and profuse pubic hair and axillary hair and voice broken), advanced puberty (advanced development of genitalia, but at least one other indicator not fully mature), early puberty (early development of genitalia, and some pubic or axillary hair or voice starting to break), and infantile (infantile genitalia or early adolescent genitalia, no pubic or axillary hair and voice not broken) (Hardy, Kuh, Whincup, & Wadsworth, 2006).

Risk factors. Birth weight, recorded to the nearest quarter of a pound, was extracted from the medical records within a few weeks of delivery and converted into grams. Birth order was reported by mothers of study members (included as a continuous measure). Height and weight measured at age 7 years, and BMI, defined as $\text{weight(kg)}/\text{height(m)}^2$, were used as indicators of pre-pubertal body size. Childhood socioeconomic position (SEP) was defined using fathers' occupation classified as non-manual (professional, managerial or intermediate)

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

or manual (skilled manual, semi-skilled manual, and unskilled) when the participant was aged 11 years, or, if this was unknown, occupation at age 4 years or 15 years.

Level of physical activity at age 36 years (most active, less active or non-active) was derived from responses to questions on level of participation in leisure activities. Life course smoking status (lifelong smoker, predominantly smoker, predominantly non-smoker, never smoker) was based on reports of smoking behaviour at all contacts since age 20 years (Clennell, Kuh, Guralnik, Patel, & Mishra, 2008). Similarly, age at birth of first child was obtained from reports at all adult follow-ups. Measured waist circumference at age 36 years was considered as a marker of abdominal adiposity. Adult SEP at age 36 years was defined as own occupation, classified as manual (skilled manual, semi-skilled manual, and unskilled) and non-manual (professional, managerial or intermediate). The highest qualification achieved by cohort members by age 26 years was classified on the Burnham scale, and was grouped into five levels: degree level and equivalents (reference group), "A" level and equivalents, "O" level and equivalents, less than "O" level, and no qualifications.

Analytical strategy

Multinomial logistic regression models were fitted to examine the associations between pubertal timing and the longitudinal profiles of affective symptoms in males and females separately. Each of the three affective symptom profiles was compared with the 'absence of symptoms' profile as the reference group. Tests for trend across the pubertal timing groups were carried out. To assess whether any observed associations with adolescent-onset affective symptoms were explained by early life risk factors, childhood SEP, birth weight, birth order, BMI and height at age 7 years were added to the regression models. To assess whether any observed associations between age at puberty and adult-onset affective symptoms were explained by behavioural or social mechanisms, the model was adjusted for adult SEP, educational attainment, exercise, smoking, age at birth of first child, and waist

circumference. We carried out additional analyses using similar models to assess whether each of the three other markers of pubertal development in girls (breast development; pigmented pubic hair; axillary hair) were associated with the affective symptom profiles.

Results

A total sample of 1972 boys and 1809 girls with valid measures of pubertal status and affective symptoms was used for the analyses. Those participants excluded from the analyses because of missing data were not different in terms of pubertal status ($p = .06$ for boys, and $p = .56$ for girls) or childhood SEP ($p = .73$ for boys, and $p = .07$ for girls) compared with those included. With regard to affective symptoms, among boys, those included in the analyses did not differ from those excluded on any of the affective symptoms profiles (all $p > .05$). Among girls, those excluded because of the missing data were more likely to have adolescent and adult symptoms ($p = .03$) and less likely to have adolescent only symptoms ($p = .01$).

Descriptive data for affective symptoms and pubertal timing, as well as for risk factors, are presented in Table 1, for males and females separately. There were sex differences in the affective symptoms groups, with a greater proportion of females being allocated to one of the three groups with symptoms ($p < .001$). Among both males and females, those with the latest pubertal timing represented the smallest group, with frequency of 10.6% for infantile group in boys and 7.5% with age at menarche ≥ 15 year in girls.

The results of the association analysis between pubertal timing and affective symptoms in females are presented in Table 2. There was no evidence that those with early age at menarche (≤ 11.11 years) were more likely to have adolescent-onset affective symptoms than normal developers (13-13.11 years). Similarly, late developers (≥ 15 years) were not at higher risk for adolescent-onset affective symptoms than normal developers. Moreover, there was no clear linear trend for increasing risk of adolescent-onset affective

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

symptoms across the groups with different pubertal timing ($p = .60$ for adolescent symptoms only; $p = .33$ for adolescent and adult symptoms). Those with the latest age at menarche (≥ 15 years) were less likely to have adult-onset affective symptoms than normal developers: OR = 0.54, 95% CI: 0.31, 0.95; $p = .03$; or early developers: OR = 0.53, 95% CI: 0.29, 0.83; $p = .04$. However, there was no clear linear trend for increased risk of adult-onset affective symptoms across the five groups with different pubertal timing ($p = .21$). The association between pubertal timing and adult-onset affective symptoms was hardly altered after adjusting for the potential confounders and mediators, i.e. adult SEP, education level, smoking status, physical activity, age at birth of first child, and waist circumference (Table 3).

Additional analyses using other markers of pubertal development at age 15 years in girls did not reveal any significant effects of pubertal development on risk of affective symptoms (Supplementary Table 1). However, these analyses were likely to lack statistical power due to the small number of girls who were not showing signs of pubertal development (i.e., breast development) at age 15 years, and therefore these findings are inconclusive.

The results of association analysis between pubertal timing and affective symptoms in males are presented in Table 4. In males, in contrast with females, late pubertal timing was associated with increased risks of affective symptoms with onset in adolescence compared to early puberty, with a clear linear trend observed across the four groups with different pubertal timing: OR = 1.31, 95% CI: 1.07, 1.60; $p = .010$ (for symptoms in adolescence), and OR = 1.21, 95% CI: 1.08, 1.34; $p = .001$ (for symptoms in adolescence and adulthood). Males with the latest pubertal timing were at increased risk of adolescent-onset affective symptoms compared to those with the earliest pubertal timing: OR = 2.59, 95% CI: 1.32, 5.09 (for adolescent only symptoms) and OR = 2.10, 95% CI: 1.44, 3.06 (for symptoms in adolescence and adulthood). There was no significant association between pubertal timing and adult-onset

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

affective symptoms in males (p for trend = .35). Associations between pubertal timing and affective symptoms with onset in adolescence were somewhat attenuated in the models controlling for childhood SEP, birth weight, birth order, BMI and height at age 7 years. However, in this fully adjusted model, the association between late pubertal timing and risk for affective symptoms in adolescence and adulthood remained statistically significant: OR = 1.87, 95% CI: 1.22, 2.86; p = .004 (Table 5). In this model, among all childhood risk factors only BMI at age 7 years showed a significant association with affective symptoms in adolescence and adulthood; those with higher BMI at age 7 years had a decreased risk of affective symptoms: OR = 0.87, 95% CI: 0.79, 0.96 (per 1 kg/m² increase; Supplementary Table 2).

Discussion

The present study revealed that late pubertal timing (age at menarche of 15 years or later) was associated with decreased risk of adult-onset, but not adolescent-onset, affective symptoms, in females. This association was not mediated or confounded by adult socio-behavioural factors. In contrast, in males, late puberty was associated with increased risk of adolescent-onset affective symptoms that continue through adult life, but not with adult-onset symptoms. This association was partially explained by low pre-pubertal BMI.

Our results suggest that in males, the negative effect of late puberty on emotional health persists into adulthood, possibly due to tracking of affective symptoms from adolescence to adulthood. To the best of our knowledge, there have not been any comparable studies following up males with different pubertal timing from adolescence through adulthood. The literature on the effects of pubertal maturation on emotional health in adolescent males is very mixed (Reardon, Leen-Feldner, & Hayward, 2009). In general, our findings are consistent with the results from several studies demonstrating that late maturing

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

boys had a higher prevalence of internalizing symptoms (Angold, Costello, & Worthman, 1998; Conley & Rudolph, 2009; Graber, Lewinsohn, Seeley, & Brooks-Gunn, 1997; Laitinen-Krispijn, van der, & Verhulst, 1999). For instance, a study in a sample of Dutch adolescents found that boys who showed greater pubertal development through the study were half as likely to exhibit parent-reported anxious and depressive symptoms as late developers (Laitinen-Krispijn, van der, & Verhulst, 1999). Another more recent study demonstrated that depression was associated with less mature pubertal status and late timing (actual and perceived) in boys (Conley & Rudolph, 2009). However, a number of studies showed that early puberty in boys was associated with higher risk of emotional problems (Ge, Conger, & Elder, 2001; Kaltiala-Heino, Marttunen, Rantanen, & Rimpela, 2003). On the whole, empirical evidence for boys suggests that both early and late maturing is associated with increased risk of affective psychopathology in adolescence. The present study adds to the existing evidence that late maturing boys can represent a high risk group for persistent affective symptoms across adolescence and adulthood. It has been suggested that different individual vulnerabilities prior to puberty may operate in interaction with pubertal timing in predicting adolescent outcomes including affective psychopathology (Benoit, Lacourse, & Claes, 2013; Gaysina, et al., 2013), and this moderating hypothesis needs to be further explored in relation to life course symptoms of depression and anxiety.

We found no evidence that early pubertal maturation (measured as age at menarche) was a risk factor for adolescent-onset affective symptoms in females. This conflicts with the generally held view of earlier pubertal maturation being associated with adolescent emotional problems in girls (Reardon, Leen-Feldner, & Hayward, 2009). However, it is worth mentioning that findings for depression (Joinson, Heron, Lewis, Croudace, & Araya, 2011; McCabe, Ricciardelli, & Banfield, 2001; O'Dea & Abraham, 1999) and anxiety (Hayward et al., 1992; Huerta & Brizuela-Gamino, 2002) are not entirely consistent. For example, a

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

longitudinal study observed a tendency for increased anxiety in Spanish girls in Tanner stage 1 (least developed) at age 12 years (Canals, Marti-Henneberg, Fernandez-Ballart, Cliville, & Domenech, 1992). A study of 286 German girls (13-14 years) showed that early maturing girls had higher risk of anxiety symptoms but not depressive symptoms (Silbereisen, Kracke, Schulenberg, Maggs, & Hurrelmann, 1997). Another study failed to confirm the association between perceived pubertal timing and internalizing symptoms in 890 girls aged 14-18 years, but did report higher risk for depressive symptoms among early maturing girls (Graber, Lewinsohn, Seeley, & Brooks-Gunn, 1997). Therefore, this research question requires further exploration, also in relation to the possible factors that could confound or mediate the observed associations. For example, BMI and/or body perception might explain the relationship of puberty with adolescents' depressive symptoms (Yuan, 2007), and these traits also vary across different samples (Ge, Elder Jr, Regnerus, & Cox, 2001).

In the present study, while there was no clear trend between pubertal timing and adult-onset affective symptoms in females, we found that the group of late developers (menarche at age 15 or later) was at lower risk than the group of on-time developers (menarche at age 13 years). In line with our finding, a US community-based cohort study of more than 4000 premenopausal women (aged 36 to 44 years) showed that later age at menarche was associated with lower risk of current depression (test for trend across four age groups: $p = .03$; Harlow, Cohen, Otto, Spiegelman, & Cramer, 1999). Contrary to these findings, a study using data from the Northern Finnish 1966 birth cohort reported that women with particularly late menarche (16 years and over) had increased prevalence of depression by age 31 years (Herva, et al., 2004). Another longitudinal study found higher prevalence of lifetime depression and anxiety in young women (age 24 years) who perceived themselves as early maturing as compared with on-time developers (Graber, Seeley, Brooks-Gunn, & Lewinsohn, 2004).

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Combined with previous findings, our results show that the association between pubertal maturation and emotional problems is different for males and females. In females, the association between early age at menarche and depression in later life was not due to tracking of this association from adolescence to adulthood, as the association was observed with adult-onset rather than with adolescent-onset symptoms. Early menarche was shown to be related to higher BMI and waist circumference in adulthood in a number of studies, including the British 1946 birth cohort (Hardy, Kuh, Whincup, & Wadsworth, 2006), which were shown to be risk factors for affective disorders (Herva, et al., 2006). However, as previously reported for this cohort, those with adult-onset affective symptoms did not differ in their BMI trajectories from those without affective symptoms (Gaysina, Hotopf, et al., 2011). This may explain why adiposity did not mediate the association between late age at puberty and reduced risk of adult-onset affective symptoms in the present study.

In males, there was an association between late age at puberty and affective symptoms with adolescent onset. The associations were partly explained by childhood risk factors, particularly childhood BMI. As reported previously, adolescent-onset affective symptoms were associated with lower BMI throughout the life course in males (Gaysina, Hotopf, et al., 2011). It is widely thought that slower childhood growth is a marker of early adverse experiences (e.g., malnutrition, chronic illness) which are also associated with affective symptoms. However, the effects of early adversity on growth have been shown to be similar in boys and girls in both population-based and adoption samples (Li, Manor, & Power, 2004; Palacios, Román, & Camacho, 2011). Thus one would expect to find similar associations between lower childhood BMI and depressive symptoms in boys and girls if the association between late puberty and depressive symptoms was fully explained by early adversity.

Sex differences in the association between pubertal maturation and affective symptoms might be explained by differential effects of hormonal influences (Martel, Klump,

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Nigg, Breedlove, & Sisk, 2009). Gonadal hormones can affect behaviour and mood differently in males and females. Gonadal hormones are known to influence HPA axis reactivity, and therefore, susceptibility to depression. Testosterone tends to reduce the peak and recovery time of the hormonal stress response in males, while ovarian steroids increase these parameters in females (Carey, Deterd, de Koning, Helmerhorst, & de Kloet, 1995; McCormick, Linkroum, Sallinen, & Miller, 2002). Thus, higher levels of estrogen in women with earlier age at menarche might increase the long-term risk for affective symptoms, and lower levels of testosterone in later maturing men may lead to increased risk of affective symptoms.

Puberty is marked by changes in neuroendocrine axes, resulting in altered hormonal output. The existing literature indicates that adolescent maturation is associated with substantial changes in stress reactivity. Specifically, it was shown that pre-pubertal rats take longer than adults to terminate an endocrine response to an acute stressor suggesting that the negative-feedback mechanisms regulating the HPA axis undergo a maturational process during puberty (Romeo, 2010). Furthermore, human studies have shown that hormonal stress reactivity increases from childhood to adolescence (Stroud et al., 2009), and that 13 and 15-year olds show greater stress reactivity than 9- and 11-year olds (Gunnar, Wewerka, Frenn, Long, & Griggs, 2009). Thus, different stress reactivity in pre- and post-pubertal stages in life may explain the differences in the association between puberty and depression in adolescence and in adulthood. However it is difficult to explain why stress reactivity and risk for affective symptoms vary by age at puberty. Future studies of the effects of pubertal timing and gonadal hormones on stress reactivity across the life span have the potential to illuminate risk and protective mechanisms for affective disorders.

Strengths and limitations

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

The present study is unique in assessing the association between pubertal timing and affective symptoms across the life course and, to our knowledge, is the first to investigate the long-term impact of timing of puberty on affective symptoms in men. Importantly, pubertal maturation in boys and girls was assessed prospectively, using both maternal reports and medical examination, as age at menarche has shown to be only moderately well recalled in adulthood (Cooper et al., 2006). In this study, we were able to adjust for pre-pubertal risk factors and to investigate pathways linking age at puberty with affective symptoms in adulthood by adjusting for a range of prospectively measured socio-behavioural factors.

The present study utilised multiple measures of symptoms of depression and anxiety assessed across the 40-year period. These multiple measures were combined using a psychometric approach that allowed using different scales for the measurement of affective symptoms at different ages, and made it possible to identify groups of individuals with differing experience of depression and anxiety over the life course (Colman, Ploubidis, Wadsworth, Jones, & Croudace, 2007). Life course phenotypes derived from repeated measurements may provide more reliable definitions of mental health phenotypes since they capture temporal aspects of an individual's vulnerability to disorder, not just diagnosis at a single time point (Leoutsakos, Zandi, Bandeen-Roche, & Lyketsos, 2010). However, one disadvantage of this approach is that people were assigned to groups according to probability. It is also worth noting that phenotypes of affective symptoms, and not clinically diagnosed affective disorders, were used in the current study, and that the pattern of results might be different in samples with a clinical diagnosis. Moreover, given that the measures of affective symptoms used in the current study were unable to distinguish between symptoms of anxiety and depression, they might mask differences in associations with pubertal timing.

Missing data and drop-outs are unavoidable in long running cohort studies such as the NSHD. The latent class approach used to define the longitudinal profiles does not require for

each individual to have all outcome measures recorded, which means that the sample size for the current analyses is maximised. There were differences for affective symptoms groups in girls: those excluded because of the missing data were more likely to have adolescent onset repeated symptoms. However, importantly, there were no differences in terms of pubertal status compared with those included and those excluded from analyses. There is no reason to suspect that the missing data would have influenced the pattern of our findings.

Although some potential confounders and mediators (e.g., SEP, health behaviours) were included in the current study, residual confounding remains a possibility, and there may be other social mechanisms involved. Other factors relevant to the effect of late menarche on adult-onset affective symptoms should be considered in the future.

In conclusion, the timing of pubertal maturation could have long-term effects on emotional problems, particularly in males. Investigating plausible biological and social mechanisms underlying these associations may lead to the development of novel suitable intervention strategies to improve mental health across the life course.

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PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Table 1. Descriptive statistics for males and females in the British 1946 birth cohort; numbers and frequencies are shown unless specified

Variables	Males	Females
Affective symptoms groups:		
- No affective symptoms	1012 (51.3%)	695 (38.4%)
- Adolescent only affective symptoms	116 (5.9%)	126 (7.0%)
- Adult only affective symptoms	263 (13.3%)	289 (16.0%)
- Adolescent and adult affective symptoms	581 (29.5%)	699 (38.6%)
Pubertal timing groups:		
Boys		
- fully mature	475 (24.1%)	
- advanced puberty	594 (30.1%)	
- early puberty	694 (35.2%)	
- infantile	209 (10.6%)	
Girls (age at menarche, years)		
- ≤ 11.11		281 (15.5%)
- 12-12.11		487 (26.9%)
- 13-13.11		626 (34.6%)
- 14-14.11		226 (12.5%)
- ≥ 15		189 (7.5%)
Girls (other markers of puberty at 15):		
Breast development, yes		1782 (99%)
Profuse pigmented pubic hair, yes		996 (55.7%)
Axillary hair, yes		1576 (88.7%)
Risk factors:		
Birth weight (gr)*	3479 (519)	3320 (483)

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Birth order:

- 1 st	809 (41.0%)	734 (40.6%)
- 2 nd or 3 rd	921 (46.7%)	861 (47.6%)
- $\geq 4^{\text{th}}$	128 (12.3%)	214 (11.8%)

BMI at age 7 (kg/m²)*

15.9 (1.3)	15.7 (1.6)
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Height at age 7 (cm)*

120.5 (5.6)	119.5 (5.7)
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Childhood SEP:

- Non-manual	807 (41.5%)	705 (39.5%)
- Manual	1137 (58.5%)	1078 (60.5%)

Adult SES:

- Non-manual	724 (54.6%)	691 (65.6%)
- Manual	601 (45.4%)	363 (34.4%)

Education level:

- Degree level and equivalents	240 (12.7%)	90 (5.2%)
- "A" level and equivalents	495 (26.3%)	364 (20.8%)
- "O" level and equivalents	282 (14.9%)	422 (24.2%)
- Less than "O" level	118 (6.3%)	176 (10.1%)
- No qualifications	749 (39.8%)	835 (39.8%)

Physical activity:

- Most active	582 (43.1%)	447 (32.9%)
- Less active	346 (25.6%)	325 (23.9%)
- Not active	423 (31.3%)	588 (43.2%)

Smoking status:

- Never smoker	334 (23.9%)	449 (31.9%)
- Predominantly non-smoker	484 (34.7%)	428 (30.4%)
- Predominantly smoker	327 (23.4%)	274 (19.5%)
- Lifelong smoker	251 (18.0%)	257 (18.2%)

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Age at birth of 1st child (years)*	26.2 (5.10)	23.6 (4.2)
Waist circumference (cm)*	89.8 (9.4)	77.0 (11.7)

*Descriptive statistic: M (SD)

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PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Table 2. Associations between pubertal timing and affective symptoms in females ($n = 1752$)

Pubertal timing (age at menarche)	Affective symptoms (AS)								
	Adolescent only AS			Adult only AS			Adolescent and adult AS		
	<i>n</i>	OR	(95% CI)	<i>n</i>	OR	(95% CI)	<i>n</i>	OR	(95% CI)
≤ 11.11	17	0.77	(0.43, 1.40)	48	1.02	(0.68, 1.54)	110	1.09	(0.79, 1.51)
12-12.11	34	0.84	(0.52, 1.35)	80	0.92	(0.66, 1.31)	178	0.96	(0.73, 1.27)
13-13.11 [§]	50	1		107	1		228	1	
14-14.11	7	0.43	(0.18, 1.00)	36	1.04	(0.66, 1.63)	105	1.42	(1.01, 2.01)
≥ 15	18	1.16	(0.64, 1.10)	18	0.54	(0.31, 0.95)	78	1.09	(0.76, 1.58)

[§]Reference group

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Table 3. Associations between pubertal timing and adult-onset affective symptoms in females: controlling for adulthood risk factors (the sample includes those with complete data; $n = 1167$ (total))

Pubertal timing (age at menarche)	Model 1: Unadjusted			Model 2: Fully adjusted*	
	<i>n</i>	OR	(95% CI)	OR	(95% CI)
≤ 11.11	40	1.18	(0.74, 1.90)	1.24	(0.76, 2.04)
12-12.11	69	1.13	(0.76, 1.68)	1.12	(0.74, 1.69)
13-13.11 [§]	85	1		1	
14-14.11	27	1.03	(0.59, 1.77)	0.99	(0.57, 1.73)
≥ 15	13	0.54	(0.27, 1.05)	0.55	(0.28, 1.08)

[§]Reference group

*Model adjusted for adult SEP, education, smoking status, level of physical activity, age at birth of 1st child, and waist circumference

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Table 4. Associations between pubertal timing and affective symptoms in males ($n = 1972$)

Pubertal timing	Affective symptoms (AS)								
	Adolescent only AS			Adult only AS			Adolescent and adult AS		
	<i>n</i>	OR	(95% CI)	<i>n</i>	OR	(95% CI)	<i>n</i>	OR	(95% CI)
fully mature [§]	21	1		82	1		112	1	
advanced puberty	33	1.32	(0.75, 2.35)	72	0.74	(0.52, 1.06)	181	1.36	(1.02, 1.82)
early puberty	44	1.52	(0.88, 2.62)	82	0.73	(0.51, 1.03)	210	1.36	(1.34, 1.80)
infantile	18	2.59	(1.32, 5.09)	27	0.99	(0.60, 1.64)	78	2.10	(1.44, 3.06)

[§]Reference group

PUBERTY AND LIFE COURSE AFFECTIVE SYMPTOMS

Table 5. Associations between pubertal timing and adolescent-onset affective symptoms in males: controlling for childhood risk factors (the sample includes those with complete data; $n = 1709$ (total))

Pubertal timing	Model 1: Unadjusted			Model 2: Fully adjusted*	
	<i>n</i>	OR	(95% CI)	OR	(95% CI)
Adolescent and adult AS					
fully mature [§]	95	1		1	
advanced puberty	156	1.40	(1.03, 1.91)	1.35	(0.99, 1.85)
early puberty	178	1.38	(1.02, 1.87)	1.31	(0.96, 1.79)
infantile	67	2.08	(1.39, 3.12)	1.87	(1.22, 2.86)
Adolescent only AS					
fully mature [§]	19	1		1	
advanced puberty	29	1.30	(0.71, 2.38)	1.30	(0.70, 2.39)
early puberty	37	1.43	(0.80, 2.56)	1.43	(0.79, 2.60)
infantile	13	2.02	(0.95, 4.28)	1.98	(0.90, 4.34)

[§]Reference group

*Model adjusted for childhood SEP, birth weight, birth order, height and BMI at age 7